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without hearing informally suspend such a permit upon the grounds set forth in paragraph (a) of this section, pending determination of formal proceedings under part 123 of this subchapter for suspension or revocation of the permit.

[23 FR 10065, Dec. 23, 1958. Redesignated at 31 FR 81, Jan. 5, 1966, and amended at 57 FR 30899, July 13, 1992]

PART 123—RULES OF PRACTICE **PROCEEDINGS** GOVERNING UNDER THE VIRUS-SERUM-TOXIN ACT

AUTHORITY: 7 U.S.C. 8301-8317; 21 U.S.C. 151-159; 7 CFR 2.22, 2.80, and 371.4.

§ 123.1 Scope and applicability of rules of practice.

The Uniform Rules of Practice for the Department of Agriculture promulgated in subpart H of part 1, subtitle A, title 7, Code of Federal Regulations, are the Rules of Practice applicable to administrative adjudicatory, ceedings under the Virus-Serum-Toxin

[42 FR 10960, Feb. 25, 1977]

PART 124—PATENT TERM **RESTORATION**

Subpart A—General Provisions

Sec.

Scope. 124.1

124.2 Definitions.

Subpart B—Eligibility Assistance

124.10 APHIS liaison with PTO.

Subpart C-Regulatory Review Period

124.20 Patent term extension calculation.

124.21 Regulatory review period determina-

124.22 Revision of regulatory review period determination.

124.23 Final action on regulatory review period determination.

Subpart D—Due Diligence Petitions

124.30 Filing, format, and content of petitions.

124.31 Applicant response to petition. 124.32 APHIS action on petition.

124.33 Standard of due diligence.

Subpart E—Due Diligence Hearing

124.40 Request for hearing.

124.41 Notice of hearing.

124.42 Hearing procedure.

124.43 Administrative decision.

AUTHORITY: 35 U.S.C. 156; 7 CFR 2.22, 2.80,

SOURCE: 58 FR 11369, Feb. 25, 1993, unless otherwise noted.

Subpart A—General Provisions

§124.1 Scope.

- (a) This parts sets forth procedures and requirements for APHIS review of applications for the extension of the term of certain patents for veterinary biological products pursuant to 35 U.S.C. 156—Extension of patent term. Responsibilities of APHIS include:
- (1) Assisting PTO in determining eligibility for patent term restoration;
- (2) Determining the length of a product's regulatory review period;
- (3) If petitioned, reviewing and ruling on due diligence challenges to APHIS's regulatory review period determinations; and
- (4) Conducting hearings to review initial APHIS findings on due diligence challenges.
- (b) The regulations in this part are designed to be used in conjunction with regulations issued by PTO concerning patent term extension which may be found at 37 CFR 1.710 through 1.791.

[58 FR 11369, Feb. 25, 1993, as amended at 64 FR 43045, Aug. 9, 1999]

§ 124.2 Definitions.

Animal and Plant Health Inspection Service (APHIS). The agency in the Department of Agriculture responsible for licensing veterinary biological products under the Virus-Serum-Toxin Act.

Applicant. Any person who submits an application or an amendment or supplement to an application under 35 U.S.C. 156 seeking extension of the term of a patent.

Due diligence petition. A petition submitted under §124.30 of this part.

Informal hearing. A hearing that is not subject to the provisions of 5 U.S.C. 554, 556, and 557 and that is conducted as provided in 21 U.S.C. 321(x).